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=> S G-CSF (L) (Treatm? OR Therapy)(5A)(renal(2A)((disease OR Failure) OR Nephropathy)) AND pd<=20041014

1 FILES SEARCHED...

2 FILES SEARCHED...

L1 6 G-CSF (L) (TREATM? OR THERAPY) (5A) (RENAL(2A) ((DISEASE OR FAILURE) OR NEPHROPATHY)) AND PD<=20041014

=> Dup Rem 11

PROCESSING COMPLETED FOR L1

L2 4 DUP REM L1 (2 DUPLICATES REMOVED)

ANSWER '1' FROM FILE MEDLINE ANSWER '2' FROM FILE BIOSIS ANSWER '3' FROM FILE CAPLUS ANSWER '4' FROM FILE EMBASE

=> D IBib ABS 12 1-4

L2 ANSWER 1 OF 4 MEDLINE on STN DUPLICATE 1

ACCESSION NUMBER: 2000388203 MEDLINE DOCUMENT NUMBER: PubMed ID: 10914557

TITLE: Safety of autologous hematopoietic stem cell

transplantation in patients with multiple myeloma and

chronic renal failure.

AUTHOR: Tosi P; Zamagni E; Ronconi S; Benni M; Motta M R; Rizzi S;

Tura S; Cavo M

CORPORATE SOURCE: Institute of Hematology and Medical Oncology, Seragnoli

University of Bologna, Italy.

SOURCE: Leukemia: official journal of the Leukemia Society of

America, Leukemia Research Fund, U.K, (2000 Jul)

Vol. 14, No. 7, pp. 1310-3.

Journal code: 8704895. ISSN: 0887-6924.

PUB. COUNTRY: ENGLAND: United Kingdom

DOCUMENT TYPE: Journal; Article; (JOURNAL ARTICLE) (RESEARCH SUPPORT, NON-U.S. GOV'T)

LANGUAGE: English

FILE SEGMENT: Priority Journals

ENTRY MONTH: 200008

ENTRY DATE: Entered STN: 18 Aug 2000

Last Updated on STN: 18 Aug 2000 Entered Medline: 10 Aug 2000

Patients with multiple myeloma (MM) and chronic renal failure have AB generally been excluded from myeloablative therapy programs followed by hematopoietic stem cell support because of the potential increase in transplant-related morbidity and mortality. We here report our experience treating six MM patients with moderate to severe renal insufficiency, with autologous stem cell transplantation. One of these patients required chronic hemodialysis since the diagnosis of MM was made. Peripheral blood stem cell collection was performed with either cyclophosphamide $5.5-7~\mathrm{g/m2}$ + G-CSF, 5 microg/kg/day (patients 1-3, 5 and 6) or G-CSF, 15 microg/kg/day alone (patient Number 4). Four patients (Nos 1-4) received autotransplant as front-line therapy, while the last two patients were treated in relapse, which occurred following prior autologous stem cell transplantation in support of melphalan, 200 mg/m2 (Number 5) or maintainance therapy with alpha-interferon (Number 6). High-dose chemotherapy administered as preparation to transplant included busulfan 12 mg/kg + melphalan 80 mg/m2 (patients 1-3 and 6) or melphalan 80 mg/m2 alone (patients 4 and 5) in order to reduce mucosal damage. Following transplant, prompt and sustained recovery of hematopoiesis was documented in all the patients; 500 PMN/microI and 20000 platelets/microI were reached after a median of 13 and 14 days, respectively. None of the patients suffered from WHO grade 3-4 infectious complications. Transplant-related toxicity included grade 3-4 oral mucositis (patients 1, 4 and 5) and veno-occlusive disease (patient Number 3). Renal function either improved or remained stable throughout the transplant period. All the patients but one responded to therapy, three of them are progression free after 2, 15 and 26 months; two relapsed after 16 and 4 months and one died from cholangiocarcinoma 7 months after transplant, while still in remission. Although our experience is limited so far, these results appear promising and support the investigational use of myeloablative therapy in MM patients with chronic renal failure.

L2 ANSWER 2 OF 4 BIOSIS COPYRIGHT (c) 2007 The Thomson Corporation on STN

ACCESSION NUMBER: 2002:186929 BIOSIS DOCUMENT NUMBER: PREV200200186929

TITLE: The Royal Marsden Hospital leukemia-myeloma database: An

"operations research" resource for assessing clinical

outcomes and planning new drug trials.

AUTHOR(S): Powles, Ray [Reprint author]; Milan, Sarah [Reprint

author]; Horton, Clive [Reprint author]; Sirohi, Bhawna
[Reprint author]; Treleaven, Jennie [Reprint author];
Singhal, Seema [Reprint author]; Mehta, Jayesh [Reprint

author]

CORPORATE SOURCE: Royal Marsden Hospital, Surrey, UK

SOURCE: Blood, (November 16, 2001) Vol. 98, No. 11 Part

1, pp. 426a. print.

Meeting Info.: 43rd Annual Meeting of the American Society of Hematology, Part 1. Orlando, Florida, USA. December

07-11, 2001. American Society of Hematology.

CODEN: BLOOAW. ISSN: 0006-4971.

DOCUMENT TYPE: Conference; (Meeting)

Conference; Abstract; (Meeting Abstract)

Conference; (Meeting Poster)

LANGUAGE: English

ENTRY DATE: Entered STN: 13 Mar 2002

Last Updated on STN: 13 Mar 2002

AB Operations research is the application of scientific methods to solve complex organizational problems including improving existing systems and designing new systems well. It is widely used in industry, finance, and

education. We have been using operations research to manage leukemia and myeloma patients for over 2 decades. Since 1978, comprehensive data have been collected prospectively on 3500+ consecutive unselected population-based patients referred to the RMH Leukaemia and Myeloma Units. For each patient, up to 600 separate data fields are recorded with multiple, longitudinal prospective values for each (e.g. serial blood counts recorded on all the days they are done - a "sparse array") on all aspects of disease and therapy. The database comprises a definition, collection and query/analysis package written in MUMPS, and presently contains apprx2X107 data items. The purpose behind its depth and breadth is to use the information for optimal clinical decision-making, outcomes analysis and planning studies on a regular basis - which also assures its quality and accuracy in an on-going fashion (Koerner 1964, Hiller-Lieberman 1974). It is possible to obtain information on a defined patient population at any instant in time (snapshot) or over a given period (panorama). For e.g., some of the information provided by a myeloma snapshot is: (a) number of living patients, (b) number in CR or PR, (c) number on maintenance interferon therapy, (d) number on induction/reinduction, (e) number with renal failure, (f) number on erythropoietin or G-CSF, etc. Some of the information provided by a myeloma panorama for the last year is: (a) number of new/relapsed patients seen, (b) number not responding to initial therapy, (c) number of patients relapsing after the first autograft, (d) number of patients failing post-autograft salvage therapy, (e) number presenting with renal failure, (f) number autografted, etc. For each group identified, detailed clinical, hematologic/biochemical data are available. Extrapolating this to the future, with specific inclusion/exclusion criteria, it is possible to predict accurately how many patients can be recruited for a study over a period of time. Thus, for a proposed 1-year study of a new drug in myeloma patients relapsing after the first autograft with normal renal function, 83 patients can be enrolled almost immediately (the number alive today with disease relapsing after first autograft) and 21 more over the next year (based upon the number relapsing over last year). By changing the inclusion to any relapse, the total number increases to 112, and by eliminating the renal function criterion, to 114. Similar analyses can also be done for leukemia. The strength of this analysis is that it can be extrapolated to other centers. It is becoming increasingly difficult to test drugs in modern health care environment due to appropriately stringent demands from drug regulatory bodies and organizations overseeing the interests of research subjects. Delays in completing pivotal trials can place large financial burdens on pharmaceutical companies which can partly be avoided by projecting enrollment based on such databases. Multi-institutional databases have larger patient numbers but suffer from several drawbacks: not having a population base (unselected consecutive patients), not being prospective, not having serial clinical data, and often having only transplant data (IBMTR, EBMT) which offer limited glimpse of the course of disease.

L2 ANSWER 3 OF 4 CAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 2002:824419 CAPLUS

DOCUMENT NUMBER: 137:304836

TITLE: Clinical application of hematopoietic factors

AUTHOR(S): Bessho, Masami

CORPORATE SOURCE: 1st Dep. Intern. Med., Saitama Med. Sch., Japan

SOURCE: Nippon Naika Gakkai Zasshi (2002), 91(Supplement, September), 229-235 CODEN: NNGAAS; ISSN: 0021-5384

PUBLISHER: Nippon Naika Gakkai DOCUMENT TYPE: Journal; General Review

LANGUAGE: Japanese

AB A review on (1) biosynthesis, distribution, and functions of

erythropoietin (EPO) and its receptor, (2) clin. application of EPO in the treatment of anemia of chronic renal failure,

(3) novel applications of EPO for the patients with cancers, myelodysplastic syndrome, or nervous system diseases, (4) clin.

applications of G-CSF in patients with neutropenia, (5) possible involvement of G-CSF administration in

the clonal complications in patients with aplastic anemia, (6)

hematopoietic growth factors for platelet production, and (7) hematopoietic disorders induced by the antibodies to hematopoietic factors (EPO, thrombopoietin, etc.).

L2 ANSWER 4 OF 4 EMBASE COPYRIGHT (c) 2007 Elsevier B.V. All rights

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ACCESSION NUMBER: 94295827 EMBASE

DOCUMENT NUMBER: 1994295827

TITLE: [The biological response modifiers. Current use and future

in clinical practice].

LES MODIFICATEURS DE LA REPONSE BIOLOGIQUE. UTILISATION

ACTUELLE ET AVENIR EN PRATIQUE LIBERALE.

AUTHOR: Kamioner D.

CORPORATE SOURCE: Dept. Hematologie/Oncologie Medicale, Centre

Medico-Chirurgical, Rue Castiglione del Lago, 78190 Trappes,

France

SOURCE: Bulletin de la Societe Française de Cancerologie Privee, (

1994) Vol. 13, No. 37, pp. 64-69. .

ISSN: 0753-7417 CODEN: BFCPE5

COUNTRY: France

DOCUMENT TYPE: Journal; Conference Article

FILE SEGMENT: 016 Cancer

037 Drug Literature Index 038 Adverse Reactions Titles

LANGUAGE: French

SUMMARY LANGUAGE: English; French

ENTRY DATE: Entered STN: 2 Nov 1994

Last Updated on STN: 2 Nov 1994

AB The development of new cytokines or biological response modifiers allows great expectations in the treatment of various tumoral diseases such as renal carcinomas, malignant melanomas, acute leukemias or infectious diseases like viral hepatitis (Interferons, Interleukines), septic shock (Centoxin), but also in hematopoietic growth disorders (G-CSF, GM-CSF, erythropoietin). However the use of these drugs is restricted because of their high cost.

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=> S CSF (L) (TReatment (3A) Nephropathy) AND pd<=20041014 2 FILES SEARCHED...

0 CSF (L) (TREATMENT (3A) NEPHROPATHY) AND PD<=20041014

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- NEWS 15 MAR 06 INPADOCDB and INPAFAMDB enhanced with new display formats
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- NEWS 17 MAR 11 ESBIOBASE reloaded and enhanced
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- NEWS 21 APR 03 CAS coverage of exemplified prophetic substances enhanced
- NEWS 22 APR 07 STN is raising the limits on saved answers
- NEWS 23 APR 24 CA/Caplus now has more comprehensive patent assignee information
- NEWS 24 APR 26 USPATFULL and USPAT2 enhanced with patent assignment/reassignment information
- NEWS 25 APR 28 CAS patent authority coverage expanded
- NEWS 26 APR 28 ENCOMPLIT/ENCOMPLIT2 search fields enhanced
- NEWS 27 APR 28 Limits doubled for structure searching in CAS REGISTRY
- NEWS 28 MAY 08 STN Express, Version 8.4, now available
- NEWS 29 MAY 11 STN on the Web enhanced
- NEWS 30 MAY 11 BEILSTEIN substance information now available on STN Easy
- NEWS 31 MAY 14 DGENE, PCTGEN and USGENE enhanced with increased limits for exact sequence match searches and introduction of free HIT display format
- NEWS 32 MAY 15 INPADOCDB and INPAFAMDB enhanced with Chinese legal status data

NEWS EXPRESS JUNE 27 08 CURRENT WINDOWS VERSION IS V8.3, AND CURRENT DISCOVER FILE IS DATED 06 APRIL 2009.

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- => S ((Kataoka, M)/au OR (Yorozu, K)/au) AND G-CSF L1 0 ((KATAOKA, M)/AU OR (YOROZU, K)/AU) AND G-CSF
- => S ((Kataoka, M)/au OR (Yorozu, K)/au) AND (Kidney OR Nephropathy)
 L2 28 ((KATAOKA, M)/AU OR (YOROZU, K)/AU) AND (KIDNEY OR NEPHROPATHY)
- => Dup Rem L2

PROCESSING COMPLETED FOR L2

L3

22 DUP REM L2 (6 DUPLICATES REMOVED)

ANSWERS '1-12' FROM FILE MEDLINE

ANSWERS '13-15' FROM FILE BIOSIS

ANSWERS '16-18' FROM FILE CAPLUS

ANSWERS '19-22' FROM FILE EMBASE

- => D L3 Ti 1-22
- L3 ANSWER 1 OF 22 MEDLINE on STN DUPLICATE 1
- TI Effects of MRX-III on chronic renal failure in rats.
- L3 ANSWER 2 OF 22 MEDLINE on STN DUPLICATE 2
- TI Effects of total parenteral nutrition containing essential trace elements on their concentrations in rats.
- L3 ANSWER 3 OF 22 MEDLINE on STN DUPLICATE 3
- TI Changes in trace-element concentrations after intravenous injection of an essential trace-element preparation for parenteral use in rats.
- L3 ANSWER 4 OF 22 MEDLINE on STN DUPLICATE 4
- TI Supplementation of essential trace elements during total parenteral nutrition—effects on trace element—deficient rats.
- L3 ANSWER 5 OF 22 MEDLINE on STN
- TI Distribution of radiolabeled cis-diamminedichloroplatinum (II) on scintigraphic images in the tumor-bearing rabbit.
- L3 ANSWER 6 OF 22 MEDLINE on STN
- TI Treatment response on CT scan in patients with brain metastases treated with irradiation.
- L3 ANSWER 7 OF 22 MEDLINE on STN
- TI A microtiter method for long-term in vitro maintenance of human monocyte.
- L3 ANSWER 8 OF 22 MEDLINE on STN
- TI Effects of a new amino acids granule on rats with ischemic chronic renal failure (author's transl).
- L3 ANSWER 9 OF 22 MEDLINE on STN
- TI Studies of AMI-U-II, a new amino acids solution for renal failure. (III)

- Effects of AMI-U-II on rabbits with ischemic chronic renal failure (author's transl).
- L3 ANSWER 10 OF 22 MEDLINE on STN
- TI Studies of AMI-U-II, a new amino acids solution for renal failure. (II) Effects of AMI-U-II on the incorporation of 15N (15N-urea) into serum albumin in the rabbit with ischemic acute renal failure (author's transl).
- L3 ANSWER 11 OF 22 MEDLINE on STN
- TI Studies of AMI-U-H, a new amino acids solution for renal failure. (I) Effects of AMI-U-II in rabbits with ischemic acute renal failure (author's transl).
- L3 ANSWER 12 OF 22 MEDLINE on STN
- TI In vitro production of pyrrolidine from proline in the presence of rat brain tissue.
- L3 ANSWER 13 OF 22 BIOSIS COPYRIGHT (c) 2009 The Thomson Corporation on STN DUPLICATE 5
- TI STUDIES ON THE TRANSFER AND DISTRIBUTION OF CARBON-14 POLY CHLORINATED BI PHENYLS FROM MATERNAL TO FETAL AND SUCKLING RATS.
- L3 ANSWER 14 OF 22 BIOSIS COPYRIGHT (c) 2009 The Thomson Corporation on STN
- TI ARSENIC DISTRIBUTION IN RATS ADMINISTERED ARSENIC-74 SUB CUTANEOUSLY AND EFFECTS OF ANTIDOTES AGAINST ARSENICAL ON THIS DISTRIBUTION.
- L3 ANSWER 15 OF 22 BIOSIS COPYRIGHT (c) 2009 The Thomson Corporation on STN
- TI IN-VIVO PRODUCTION OF PIPERIDINE FROM PIPECOLIC-ACID IN THE RAT.
- L3 ANSWER 16 OF 22 CAPLUS COPYRIGHT 2009 ACS on STN
- II Influence of Zn, Cu, Mn-mixture solution on metallothionein concentration and SOD activity of organ in rats fed Zn, Cu, Mn-deficient diet
- L3 ANSWER 17 OF 22 CAPLUS COPYRIGHT 2009 ACS on STN
- TI Experimental study on intralymphatic chemotherapy using 195mPt-cis-diamminedichloroplatinum(II) with high specific activity
- L3 ANSWER 18 OF 22 CAPLUS COPYRIGHT 2009 ACS on STN
- TI Influence of an essential trace elements agent for parenteral use on metallothionein concentration in mice
- L3 ANSWER 19 OF 22 EMBASE COPYRIGHT (c) 2009 Elsevier B.V. All rights reserved on STN
- TI Thermoradiotherapy combined with daily administration of low dose cisplatin for locally advanced laryngeal carcinoma.
- L3 ANSWER 20 OF 22 EMBASE COPYRIGHT (c) 2009 Elsevier B.V. All rights reserved on STN
- TI Successful chemoradiotherapy for locally advanced gastric non-Hodgkin's lymphoma in a 3-year-2-month-old girl.
- L3 ANSWER 21 OF 22 EMBASE COPYRIGHT (c) 2009 Elsevier B.V. All rights reserved on STN
- TI A new radioiodinated ligand for SPECT studies of peripheral benzodiazepine binding sites: 4'-Iodo-PK11195.
- L3 ANSWER 22 OF 22 EMBASE COPYRIGHT (c) 2009 Elsevier B.V. All rights reserved on STN
- TI General pharmacological studies of MRX-III, a new amino acid solution for renal failure.

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